

SHARKOV, V.I.; KUYBINA, N.I.; SOLOV'YEVA, Yu.P.

Extraction of hemicelluloses from woodpulp without cleavage
of the acetyl groups. Zhur. prikl.khim. 33 no.11:2571-2575
N '60. (MIRA 14:4)

1. Gosudarstvennyy nauchno-issledovatel'skiy institut gidroliznoy
i sul'fitno-spirtovoy promyslennosti.
(Hemicellulose)

TROYANSKIY, Ye.A.; SOLOV'YEVA, Yu.P.

Nature of the yield strength area. Fiz. met. i metalloved. 12
no.5:758-759 N '61. (MGRA 14:12)

1. Moskovskiy energeticheskiy institut.
(Steel--Testing)

SHAKOV, V.I.; KUYBINA, N.I.; SOLOV'YEVA, Yu.P.; GVOZDEVA, Z.N.; ARTEM'YEVA, I.S.

Chemical composition of the corncob. Gidroliz. i lesokhim.prom.
15 no.2:7-8 '62. (MIRA 18:3)

1. Gosudarstvennyy nauchno-issledovatel'skiy institut gidroliznoy
i sul'fitno-spirtovoy promyshlennosti.

TROYANSKIY, Ye.A., kand. tekhn. nauk; KORIKOVSKIY, I.K., inzh.;
SOLOV'YEVA, Yu.P., inzh.

Methods for approximate calculation of welded T-pipes.
Teploenergetika 10 no.8:43-46 Ag '63. (MIRA 16:8)

1. Moskovskiy energeticheskiy institut.
(Steampipes—Welding)

TRIGANOV, Ye.A.; SOLODOVNIKOVA, Yu.P.

Propagation of plastic deformation. Metallized. i term. obr. met.
no.7:II-13 JI '64. (MIRA 17:01)

i. Moscowvskiy energeticheskiy institut.

SHAROV, V.I.; KIVIRINA, N.I.; S. LEV'YAK, Yu.I.

Composition and structure of zylezonide obtained by alkaline extraction. Zhur. prikl. khim. 37 no.12:2720-2725 D '64.
(MIEA 18:?)

CHIRKOV, V. I., AGBIBIN, R. I., KALUGOV, V. N. P.

Study of the homogeneity of xylcurealde by fractionation. Zhur. prikl. khim. 38 no. 7:1595-1598 JI '65. (M721 1812)

1. Gosudarstvennyy nauchno-issledovatel'skiy institut gidroizotnoy i sull'fatsospirtovoy promyshlennosti.

ALESKOVSKIY, A.M.; SOLOV'YEV, Yu.V.; ANDRUSHKEVICH, V.S.

Magnetic compensating manometers. Prib. i tekhn. eksp. no.1:110-112
Ja-P '57. (MLRA 10:6)

1. Saratovskiy gosudarstvennyy universitet im. N.O. Chernyshevskogo.
(Manometer)

GOLOVEVA, Yu.V. and ANANEVA, E.P.

"Etiology of Toxic Diphtheria," Zhu. MEIB, V. 1, No. 2, pp. 41-45,
1945.

Bact. Inst. Khar'kov.

DANSKER, V.N., RUBEL', S.N.; SOLOV'YEVA, Ye.V.

Materials on the study of leptospiral jaundice. Report no. 9: Parallelism in the laboratory and clinical aspects of leptospiral and nonleptospiral jaundice. Trudy Len.inst.epid. i mikrobiol. 9:65-83 '47.

(MILB. 10:9)

1. Iz laboratorii po izucheniyu leptosirozov (zav. K.N.Tokarevich)
Instituta epidemiologii i mikrobiologii im. Petersa (dir. P.I.Krasnits)
i infektsionnogo otdeleniya bol'ničay im. Very Slutskoy (glavnyy vrach
E.M.Abkin)

(LENINGRAD--HEPATITIS, INFECTIOUS)

(LENINGRAD--WEIL'S DISEASE)

ZIL'BER, L.A.; SOLOV'YEVA, Yu. V.; VOLINA, E.V.; KRAVCHENKO, N.A.

Antibacterial action of hemin and its derivatives. Biokhimiya 18,
109-11 '53.
(MLRA 6:1)
(CA 47 no.15:7594 '53)

1. Central Inst. Epidemiol. Microbiol., Moscow.

SOLOV'YEVA, Yu. V.

"Material Precipitated From S. Diphtheriae Which Possess Antibacterial Characteristics." Proceedings of Inst. Epidem and Microbiol im. Gamaleya 1954 56,

List of Works Sponsored by the institute [Authors are not identified with any specific division, laboratory, or other organizational component of the institute.] Inst. Epidem and Microbiol im. Gamaleya AMS USSR.

SO: Sum 1186, 11 Jan 57.

SOLOV'YEVA, Yu. V.
USSR/Microbiology - Antibiosis and Symbiosis, Antibiotics.

F-2

Abs Jour : Ref Zhur - Biol., No 4, 1958, 14718

Author : Solov'eva, Yu.V., Belova, Z.N.
Inst : -
Title : Mycerin.

Orig Pub : V sb.: Antibiotiki, Eksperim.-klinich. izuch. M., 1956,
207-213

Abstract : A new antibiotic, mycerin, represents a polypeptide and contains reducing substances in its molecule. It does not lose its activity in the presence of serum, possesses a wide spectrum of antibacterial activity, and low toxicity (150 mg/kg in intramuscular injections and 1.5 g in oral application). It exerts no cumulative effect (when introduced 100 mg/kg for a period of 5 days). After intramuscular injection in doses of 200 and 100 mg/kg this antibiotic is found after 24 hours in blood serum, while after the first 6 hours its concentration is respectively

Card 1/2

SOLOV'YEEVA, Yu.V.

Substance isolated from *Corynebacterium diphtheriae* possessing
antibacterial properties. Zhur. mikrobiol. epid. i immun. 27 no.2:
41-42 F '56. (MLRA 9:5)

1. Iz Instituta epidemiologii i mikrobiologii imeni N.F. Gamalei
AMN SSSR.
(*CORYNEBACTERIUM DIPHTHERIAE*) (BACTERICIDES)

GOLOV'YEVA, Yu. V. (Cand. of Med. Sci.); BELOVA, Z.H.;

"Mycerin (Mitserin),"

p. 207 Ministry of Health USSR Proceedings of the Second All-Union Conference on
Antibiotics, 31 May - 9 June 1957. p. 405, Moscow, Medgiz, 1957.

SOLOV'YEVA, YU. V., EREKT, M. K.

"Effect of the new antibiotic, aurantin, on the development and
recurrent growth of M-1 sarcoma."

report submitted at the 13th All-Union Congress of Hygienists, Epidemiologists
and Infectionists, 1959.

PLANEL'YES, Kh.Kh., prof., red.; SOLOV'YEVA, Yu.V., kand. med. nauk, red.; GRACHEVA, N.P., kand. med. nauk, red.; ANTONOV, B.N., red.; ZUYEVA, N.K., tekhn. red.

[Mycerin; experimental study and the results of its clinical testing] Nitserin; eksperimental'noe izuchenie i rezul'taty klinicheskogo ispytaniia. Moskva, Medgiz, 1961. 307 p.
(MIRA 15:3)

1. Akademiya meditsinskikh nauk SSSR, Moscow. Institut epidemiologii i mikrobiologii. 2. Chlen-korrespondent Akademii meditsinskikh nauk SSSR (for Planel'yes).
(ANTIBIOTICS)

PLANEL'YES, Kh.Kh., prof., red.; SOLOV'YEVA, Yu.V., kand. med. nauk, red.; GRACHEVA, N.P., kand. med. nauk, red.; ANTONOV, B.N., red.; PARAKHINA, N.L., tekhn. red.

[Aurantin, an antineoplastic antibiotic ; reparation from the actinomycin group; experimental study and the results of a clinical test]Aurantin - protivoopukholevyi antibioticheskii preparat iz gruppy aktinomitsinov; eksperimental'noe izuchenie i rezul'taty klinicheskogo ispytaniia. Moskva, Medgiz, 1962. 279 p. (MIRA 15:9)

1. Akademiya meditsinskikh nauk SSSR, Moscow. 2. Chlen-korrespondent Akademii meditsinskikh nauk SSSR (for Planel'yes).
(CYTOTOXIC DRUGS) (ANTIBIOTICS)

ORLOV, V.; SOLOV'YEVA, Z.; RUDNOVA, A., inzhener-khimik; KOVALEV, N.;
KHAKHEL', L.

Draw ship repair plant laboratories into doing creative work.
(MIRA 15:12)
Mor. flot 22 no.11:36-37 N '62.

1. Nachal'nik TSentral'noy laboratorii Rizhskogo sudoremontnogo zavoda (for Orlov).
2. Starshiy inzhener-fizik TSentral'noy laboratorii Rizhskogo sudoremontnogo zavoda (for Solov'yeva).
3. Starshiy tekhnik TSentral'noy laboratorii Rizhskogo sudoremontnogo zavoda (for Kovalev).
4. Starshiy laborant TSentral'noy laboratorii Rizhskogo sudoremontnogo zavoda (for Khakhel').

(Ships—Maintenance and repair)

SOLOV'YEVA, Z. A.

Characteristics of cerebral neurodynamics in hypertension with
a tendency to repeated critical states. Nauch. trudy Inst. nevr.
AMN SSSR no.1:210-217 '60. (MIRA 15:7)

1. Institut nevrologii AMN SSSR.

(HYPERTENSION) (CEREBRAL CORTEX)
(CEREBROVASCULAR DISEASE)

YARULLIN, Kh.Kh.; SOLOV'YEVA, Z.A.

Characteristics of conditioned and unconditioned vascular reflexes in acute disorders of the cerebral circulation. Zhur. nevr. i psikh. 62 no.1:51-58 '62. (MIRA 15:4)

1. Institut nevrologii (dir. - prof. N.V.Konovalov) AMN SSSR, Moskva.
(CEREBROVASCULAR DISEASES) (CONDITIONED RESPONSE)
(REFLEXES)

YARUILLIN, Kh.Kh.; SOLOV'YEVA, Z.A.

Physiological characteristics of disorders of consciousness in
the acute period of a cerebral insultus. Zhur. nevr. i psich.
64 no.10:1506-1514. '64. (MIRA 17:11)

1. Institut nevrologii (direktor - prof. N.V. Konovalov) AMN
SSSR, Moskva.

MOISEYENKO, U.I.; SOLOV'YEVA, Z.A.; KUTOLIN, V.A.

Heat conductivity of granite in the case of high temperature.
Dokl. AN SSSR 165 no.3:670-671 N '65. (MIRA 18:11)

1. Institut geologii i geofiziki Sibirskogo otdeleniya AN SSSR.
Submitted May 29, 1965.

4

CA

Passivation of the cathode and structure of the electrolytic deposit in solutions of simple and of complex salts. A. T. Vaginyan and Z. A. Solov'yova. *Doklady Akad. Nauk SSSR*, 68, 121-124 (1949). Ag cathodes of 0.01 cm^2 were first subjected to 5 min polarization with a dep. of 50×10^{-6} amp, then the current was turned off for t sec ($t = 0, 10, 30, 60$, and 300), and the cathode polarized again. The cathode potential φ against an equal Ag electrode was recorded beginning with the last 1 sec of the rest period, and for about 20 sec under the polarizing current. In a 0.2 N soln of AgNO_3 , polarization gives rise to a momentary upward jump of the potential to φ_{max} , followed by a fall to a steady-state φ_0 . The difference $\Delta\varphi = \varphi_{\text{max}} - \varphi_0$ increases with the length t of the preceding rest period, and characterizes the degree of passivation of the surface. Curves of $\Delta\varphi$ as a function of t are termed passivation velocity curves. Addition of trivalent dextrin increases the rate of passivation. In contrast to the simple salt, no potential jump $\Delta\varphi$ is present in the complex electrolyte (0.2 N $\text{AgNO}_3 + 0.02 \text{ M K}_3\text{Fe(CN)}_6$). After the current is turned off, the potential has a momentary value which falls slowly with time, and reaches a steady-state value, in other words, the Ag electrode in a complex electrolyte passes over an increased activity. At constant $t = 30$ sec, and constant polarizing current of 1000 amp/cm², the curves of the potential φ against the time, show, in AgNO_3 , a fall of φ ; in the presence of dextrin, the initial φ_{max} is markedly higher and the initial fall is somewhat faster. In the complex electrolyte, $\Delta\varphi$ increases with time t , at a lower dep. (1000 amp/cm²), the shape of the curve remains the same, only the φ_0 are smaller. The absence of passivation in the complex electrolyte is explained, on the basis of the work of Franklin and Oberholzer, by adsorption of the complex anions which thus prevents absorption of extraneous matter.

The smooth structure of the deposit is due to uniformity of the surface (absence of positive regions), and deposition of metal from the adsorbed ions. The dependence of $\Delta\varphi$ on solns of simple salts one refutes all theories linking the initial max. of polarization with such processes as exchange of the double layer, counter polarization or formation of crystal germs. The latter explanation is further refuted by the absence of a max. in complex electrolytes. The increase of $\Delta\varphi$ with the length t of the rest period is due, either, to a decrease of the active fraction of the surface and a corresponding increase of the true dep. This phenomenon is suppressed by the absorption of complex ions.

CA

New method of investigation of cathodic passivation
A. I. Vagruyan and Z. A. Sudor (Institute of Electrochemistry, USSR
Academy of Sciences, Kiev, Ukr. SSR 26, 1232 611 (1960); cf. t. 1
60, 400). After the const. current I of electrolysis has
been interrupted during t sec., one measures a polarization
 η at time $t = 0$, i.e. when the circuit is closed again which
must be higher than η_0 , the polarization at $t = 0$. The dif.
ferrone $\Delta\eta = \eta_0 - \eta_0$ is due to the passivation of the cathode
during t . A knowledge of the function $\Delta\eta = f(t)$ gives in
formation on the rate of passivation in various electrolytes.

and on the influence of various surface-active compounds
on the rate. An automatic app. is described in detail
it records photographically the changes of η after current
interruptions $t = 3.5, 10, 30, 60$, and 300 sec. With this
app., passivation during current interruption is observed
on a Ag cathode in a 0.2 N AgNO_3 soln, but no passivation is
found on the same cathode in a 0.2 N KAgCN soln.
Surface-active substances increase $\Delta\eta$ in the first case but
have no influence in the second. The absorption of octyl
al., on a Zn cathode was also studied. First, a polarization
curve $\eta = f(t)$ where t is the c.d. is detd. for the Zn
cathode in a 2 N $\text{ZnSO}_4 + 0.01 \text{ N NaCl}$ soln. From this
curve and from the values of η_0 detd. in the same soln
without and with octyl al., one dets. the change in η_0 ,
and thus (since $I = \text{const.}$) the decrease in active surface
 ΔS due to adsorption. Adsorption rate curves ($\Delta S, S, t$)
show that the rate is large at first and slows down when
 S_{min} is approached. Adsorption isotherms might be detd.
in this fashion. The decrease in active cathode surface
 ΔS amounts to 84.7% after $t = 30$ sec. in a soln. with
octyl al.

Michel Boudart

SOLOV'YEVA, S. A.

"Effect of Surface-Active Agents on the Process of the Electro-deposition of Metals." Sub 20 Dec 51, Inst of Physical Chemistry, Acad Sci USSR.

Dissertations presented for science and engineering degrees in Moscow during 1951.

SO: Sum. No. 480, 9 May 55

Chemical Sciences

SOLOV'YEVA, Z.A.

Effect 4

"Galvano-Polarization in the Electrodeposition of Nickel.
A. T. VASIL'YEV AND Z. A. SOLOV'YEVA (Doklady Akad. Nauk SSSR, 1957, 77, (4), 859-861).—[In Russian]. V. and Z. obtained the dependence of polarization (η) on e.d. by uniformly increasing the current (i) through the cell from zero to some max. value and then returning it to zero; this was done within a short time (16, 1.5 sec., and less), η being recorded photographically. In some experiments i was varied linearly, in others logarithmically. The cell used contained (g./l.): $\text{NiSO}_4 \cdot 6\text{H}_2\text{O}$ 140, H_3BO_3 30, KCl 19, temp. 29°C. After switching off the current, the electrode potential had a high negative value which gradually changed in the positive direction. This change also occurred in the polarization of an electrode by a small current. From curves showing changes in η for a Ni electrode when the e.d. varied linearly from 0 to 30 to 0 m.amp./cm.² the value of η at zero e.d. was —308 mV. for a 16-sec. cycle, and —212 mV. for a 1.5-sec. cycle. Taking 308 mV. as the equilibrium potential φ_0 , the overvoltage is 486 mV.; if the "steady" value (at e.d. of 30 m.amp./cm.²) be taken as φ_0 , the overvoltage is 790 mV. The curves of η/\log e.d. are linear over a greater range of e.d. for a 1.5-sec. cycle than a 16-sec. cycle, but the max. value of η is independent of speed. Glazstone's explanation of chem. polarization in terms of an active or metastable deposit (*J. Chem. Soc.*, 1926, 129, 2387; *J. Inst. Metals* (Abstracts), 1927, 87, 627) is not supported by the fact that scraping extricates the surface of the electrode; more probably a completely pure Ni surface cannot exist for a prolonged time because foreign particles are absorbed on it and cause passivation. The passive film is absent from a scraped surface or from freshly deposited Ni, and hence the potential is more negative. The change in potential of freshly deposited Ni with time to a more positive value is explained either by passivation or by the as deposited surface not having a normal equilibrium lattice, but contg. an excess of active places, and then undergoing ordering as time elapses. In any case, polarization is not wholly reduced to the slowness of the discharge on the pure surface, but indicates some change in its physico-chem. condition G. V. E. T.

VAGRAMIAN, A. T.; SOLOV'IEVA, Z. A.

Nickel

Investigation of brightness of electrolytic deposits of nickel during electrolysis.
Zhur.fiz.khim., 16, No. 6, 1952.

Monthly List of Russian Accessions, Library of Congress, November 1952. Unclassified.

Electrochemistry - 4

CP

Gloss of electrolytic nickel deposits during electrolysis
A. F. Vagrameyan and Z. A. Soker'eva (Inst. Phys. Chem.,
Acad. Sci., Moscow). *Zhur. Tekhn. Kibernetika* 20, 910-17 (1952)
A light beam was reflected from a cathode on which Ni was
being deposited, and the intensity I of the reflected beam
was measured. Thus the change of gloss caused by air was
eliminated. The cathode was of polished Fe, and the soln.
contained $\text{NiSO}_4 \cdot 7\text{H}_2\text{O}$ 140, H_3BO_4 15, NaCl 5, and NaF 1.5
 g/l . The curve " I against δ " (δ is the thickness of the Ni
deposit) had 2 parts. Between $\delta = 0$ and $\delta = 2$ or 3μ I
always increased with δ . In this region the surface profile of
Ni repeated that of Fe underneath, and the increase of I was
due to the greater reflection coeff. of Ni. At δ greater than
 3μ , I increased very little with δ (up to 10μ was investigated)
when c.d. was 1-1.5 amp./sq. dm. and decreased on an
increase of δ when c.d. was 0.3 or 2. This was explained by
the properties of the surface obtained; an electron micro-
scope showed that the deposit consisted of coarse crystals at
high c.d., of crystals of different sizes at small c.d., and of
uniform smooth crystals at medium c.d. Addns. of CS_2
 $(\text{NH}_3)_2$ (e.g., 0.15 g./l.) or of disulfonaphthalenic acid (e.g.,
4 g./l.) raised I above the highest value observed in inorg.
electrolyte and also made I almost independent of c.d. be-
tween 1 and 4 amp./sq. dm. These addns. give rise to
small uniform crystals. If the cathode was kept without
current for several sec. or min. and current was then switched
on, I decreased for several sec. This was caused by the
passivation of the electrode during the interruption of cur-
rent; when the deposition was resumed it occurred on the
spots least passivated and, thus, the cathode became less
uniform. α -Naphthol at c.d. 2 depressed I . J. J. B.

SOLOV'YEVA, Z.-A.

USSR

✓ Investigation of cathodic polarization with simultaneous
discharge of ions of iron and tungsten. Z. A. Solov'yeva
and A. T. Vayramyan. Bull. Acad. Sci. U.S.S.R., Div. II
Chem. Sci. 1954, 185-9 (Engl. translation).—See C.A.
43, 134871. H. L. H.

Soviet Union, U.S.S.R.

USSR/Chemistry Physical chemistry

Card : 1/1

Authors : Vagramyan, A. T., and Solovyeva, Z. A.

Title : Method of studying the rate of electrochemical reactions

Periodical : Zhur. fiz. khim. 28, Ed. 6, 1153 - 1157, June 1954

Abstract : A method of studying the rate of electrochemical reactions, based on complex investigation of relations between polarization and current density, between polarization and time at constant current density and between current intensity and time at constant cathode potential, is described. A more complete and accurate understanding of the processes taking place on the electrode during electro-deposition of metals can be obtained only through study of above mentioned relations. Four USSR and 2 German references. Graphs, drawing.

Institution : Acad. of Sc. USSR, Institute of Physical Chemistry, Moscow

Submitted : December 27, 1953

SOLOV'YEV, V. D.

3

*Investigation of Cathodic Polarization During Simultaneous Discharge of Iron and Tungsten Ions. Z. A. Solov'yev and A. I. Vagramyan [Izdat. Akad. Nauk S.S.R. R., 1954, (Khim.), (2), 230-233].—[In Russian]. Electrodeposition of Fe, W, and of an Fe-W alloy from citric acid soln. neutralized with NH₄OH (pH 8-8.5) was studied at 19° and 70° C. and c.d. 5 amp./dm.². The deposition of the Fe-W alloy at 19° and 70° C. took place at 978 and 880 mV., resp. The lowering of the deposition potentials with respect to deposition potentials of pure W and Fe was 320 and 109 mV. at -20° C., and 430 and 208 at 70° C. The decrease of the deposition potential during simultaneous deposition of W and Fe points to the formation of an intermetallic compound of the Fe₃W type. The absence of the periodic oscillations of the p.d. during electrodeposition of Fe-W alloy disproved the theory advanced by Holt and Vaaler (*Trans. Electrochem. Soc.*, 1948, 94, 60; see *M.A.*, 18, 293), which stated that Fe and W were deposited on the cathode in alternating layers and not simultaneously.

—S. K. L.

VAGRAMYAN, A.T.; SOLOV'YEVA, Z.A.; ALEKSEYEV, V.A., redaktor; SHIVCHENKO, G.N., tekhnicheskiy redaktor

[Research methods in the processes of electrolytic precipitation of metals] Metody issledovaniia protsessov elektroosazhdeleniya metallov. Moskva, Izd-vo Akad. nauk SSSR, 1955. 250 p. (MIRA 8:7)
(Electroplating)

Galena, Z. A.

2.7
✓ Determination of bi- and trivalent titanium in fused alkali metal chlorides after electrolysis. E. G. Tabakova and

Z. V. Golovina. Zavodskaya Lab. 22, 1417-19 (1955). *4*

The Ti-contg. melt to be tested is the electrolyte for Ti refining (a eutectic KCl + NaCl mixt., or either of the chlorides alone). A sample is dissolved in an excess of Fe NH₄ alum soln. in a stream of CO₂, and the amount of iron reduced by Ti⁴⁺ and Ti³⁺ is titrated with 0.02N KMnO₄ soln. (after adding the Zimmermann-Reinhardt soln.). Another sample is dissolved in 0.1N HCl in a CO₂ stream, adding some satd. (NH₄)₂SO₄ soln. to stabilize Ti⁴⁺, and raising the acidity of the soln. by adding 6% H₂SO₄ soln., which will oxidize Ti⁴⁺ to Ti⁴⁺, and again titrating with KMnO₄. *W. M. Sternberg.*

SOLDYKEVA, Z. A.

The effect of the acidity of a solution on the cathode polarization in the electroprecipitation of cobalt and nickel.
Z. A. Soldy'eva and O. A. Abramov (Inst. Phys. Chem., Acad. Sci. U.S.S.R., Moscow). Zhur. Fiz. Khim. 30, 1572-8(1956).—The effect of pH on the cathode polarization during the electropptn. of the Fe-group metals was studied and a new phenomenon was observed, i.e., the polarization changes sharply in a narrow pH range. This range is displaced toward lower pH values with increasing temp. This phenomenon is related with hydrate formation in solns. of Co salts, thereby indicating that this phenomenon is of a chem. nature. J. Frater-Leach

Chem

7 2

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RM
MK

SOLOV'YEVA, Z.A.

Distr: 4E2c

Effects of boric acid on the cathodic polarization during the electrodeposition of cobalt and nickel. Z. A. Solov'yeva and O. A. Abramov (Inst. Phys. Chem. Acad. Sci. U.S.S.R., Moscow, Zhur. Fiz. Khim. 31, 1248-55 (1957); cf. U.S. 3,639,746). The effect of the addn. of various amounts of H₃BO₃ upon the Co and Ni electrodeposition from CoCl₂, CuSO₄, and NiSO₄ solns. in a wide range of pH was studied voltammetrically against H₂ electrodes. H₃BO₃ raised the cathodic polarization the more the higher its concn., and the rise was higher at higher pH and lower temps. The slope of the $\eta - \log i$ polarization curve was greatly changed by the H₃BO₃ addn. The formation possibility of chem. compds. between the metal ions and H₃BO₃ was suggested, with the discharge of the metal ion on the cathode being correspondingly slowed down. W. M. Steinberg

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1

FRUMKIN, A.N., akademik, otv.red.; YESIN, O.A., prof., red.; ZHDANOV, S.I.,
red.; KABANOV, B.N., prof., red.; KOLOTYRKIN, Ya.M., dokt.khim.
nauk, red.; LOSEV, V.V., red.; LUKOVTSOV, P.D., prof., red.;
SOLOV'YEVA, Z.A., red.; STENDER, V.V., prof., red.; FLORIANOVICH,
G.M., red.; YEGOROV, N.G., red.izd-va; PRUSAKOVA, T.A., tekhn.red.

[Proceedings of the 4th Conference on Electrochemistry. October
1-6, 1956] Trudy 4-go soveshchaniia po elektrokhimii. 1-6 oktiabria
1956 g. Moskva, Izd-vo Akad.nauk SSSR, 1959. 867 p. (MIRA 12:5)

1. Soveshchaniye po elektrokhimii, 4th, Moscow, 1956.
(Electrochemistry)

SOLOV'YLOVA, Z. A.

PHASE I BOOK EXPLOITATION

SOV/4828

Vagramyan, Ashot Tigranovich, and Zoya Alekseyevna Solov'yeva

Metody issledovaniya elektroosazhdeleniya metallov (Methods of Investigating the
Electrodeposition of Metals) 2d ed., rev. and enl. Moscow, Izd-vo AN SSSR, 1960.
447 p. Errata slip inserted. 5,000 copies printed.

Sponsoring Agency: Akademiya nauk SSSR. Institut fizicheskoy khimii.

Ed. of Publishing House: N.G. Yegorov; Tech. Ed.: G. A. Astaf'yeva.

PURPOSE: This book is intended for metal scientists and persons working in the
electroplating industry.

COVERAGE: The book deals with electrochemical, physical, and mechanical research
methods used in the electrodeposition of metals from aqueous solutions. The
methods considered are primarily those used or developed at the laboratoriya
elektroosazhdeleniya metallov (Laboratory for the Electrodeposition of Metals) of
the Institut fizicheskoy khimii AN SSSR (Institute of Physical Chemistry
AS USSR). They include, among others, the method for studying electrode polarization

Card 1/10

SOLOV'YIVA, Z.A.; UVAROV, L.A.; VAGRAMYAN, A.T.

Rate of exchange between cobalt and its ions in solution. Zhur.
neorg.khim. 5 no.6:1185-1188 Je '60. (MIRA 13:7)
(Cobalt)
(Reduction, Electrolytic)
(Ion exchange)

AUTHOR:

Solov'yeva, Z. A. (Moscow)S/076/60/034/03/007/038
B115/B016

TITLE:

Investigation of the Adsorption Rate of Surface-active Substances
on the Metal SurfacePERIODICAL: Zhurnal fizicheskoy khimii, 1960, Vol 34, Nr 3, pp 537 - 542
(USSR)

TEXT: In the present paper the influence exercised by the electrolyte concentration upon the adsorption rate of saturated aliphatic alcohols in the adsorption on the surface of a zinc electrode is investigated. The study of the adsorption of octyl-, hexyl-, amyl-, and butyl alcohol in solutions of $ZnSO_4$ with different concentrations (1 - 6 N) by application of the electrochemical method showed that the dependence of the saturation of the surface on the adsorption time in a solution with alcohol added considerably depends on electrolyte concentration. The dependence of the polarization increase $\Delta\gamma$ on the time t in a 4 N $ZnSO_4$ solution + $2.5 \cdot 10^{-4}$ mole/l octyl alcohol is investigated (Fig 1), while in the following figure (Fig 2) the same dependence is given for a 1 N $ZnSO_4$ solution. It may be seen from the given diagrams that the adsorption equilibrium in the 1 N $ZnSO_4$ solution is established about six times more quickly

Card 1/3

Investigation of the Adsorption Rate of Surface-active Substances on the Metal Surface

8/076/60/034/05/007/038
B115/B016

than in the 4 N $ZnSO_4$ solution. The establishing rate of the adsorption equilibrium is not changed by a variation of the octyl alcohol concentration, but only the value $\Delta\gamma$ (Fig 3). The dependence $\Delta\gamma, \tau$ on continuous vigorous supply of the freshly prepared electrolyte to the cathode is investigated as well (Fig 4). Figure 5 shows the $\Delta\gamma, \tau$ curves in 1 and 4 N $ZnSO_4$ solutions saturated with butyl alcohol, the solubility of which is by some orders of magnitude higher than that of octyl alcohol. The effect of butyl alcohol as well as of hexyl and amyl alcohol was similar to that of octyl alcohol. Also the behavior of the salts Na_2SO_4 , $MgSO_4$, and $Al_2(SO_4)_3$ was the same as that of $ZnSO_4$. It may be said from the results obtained that the increase of the electrolyte concentration effects a change in the structure of the double layer, mainly of the packing, and that the entrance of the organic molecules into the layer is considerably inhibited. The results obtained by O. K. Gal'dikene, S. I. Khotyanovich, and Yu. Yu. Matulis (Ref 8) are mentioned. The author expresses her gratitude to A. T. Vagramyan for his assistance and advice. There are 5 figures and 8 Soviet references.

Card 2/3

SOLOV'YEVA, Z.A.; VAGRAMYAN, A.T. (Moscow)

Role of self-adjustment in polarization measurements during the
electrodeposition of metals. Zhur. fiz. khim. 34 no.4:754-758
Ap '60. (MIRA 14:5)

(Polarization (Electricity)) (Electroplating)

S 1905/080/61/034/008/007/018
D204/D305

AUTHORS Solov'yeva, Z. A. and Petrova, Yu. S.

TITLE: On the rates of combined reactions during electrodeposition of chromium

PERIODICAL: Zhurnal prikladnoy khimii, v. 34, no. 8, 1961,
1752-1759

TEXT: The object of the article was to study the effect of electrolyte composition and conditions of electrolysis on the rates of the combined reactions during electrolysis of chromic acid and to correlate the rate of each reaction with the properties of the cathode film. The relative variation of each of the combined reactions to the others was determined from the variation of current distribution between the reactions. Analytically pure CrO₃ was freed from traces of H₂O₄ by repeated washing with concentrated NaOH with subsequent drying. Potentials were measured by a compensation technique with an electrode of 1.5 cm² surface area, current distribution being simultaneously determined as between the three reactions.

Card 1/3,

On the rates of combined reactions...

5/080/61/034/008/007/018
D204/D305

An electrode of 0.002 cm² was used when the decrease of voltage per ohm of electrolyte was high. The electrode surface was Cr-plated. Experiments were made with solutions containing 250 g/l CrO₃ and varying concentrations of H₂SO₄, with temperatures of 20°, 40° and 60°, and current densities of 250, 650 and 1000 mA/cm². At the two higher temperatures, the curves depicting the effect of H₂SO₄ concentration on reaction rates show consistent dependence of the rates on acid concentration. Deposition of Cr passes through a maximum and Cr⁶⁺ → Cr³⁺ increases and 2H⁺ → H₂ decreases with increase of H₂SO₄ concentration. At current densities above 250 mA/cm², H₂SO₄ concentration retains its character in relation to reaction rates but is less marked for Cr⁶⁺ → Cr. A relation was found between the rate of electrochemical reduction of chromic acid and composition, structure and cathode film resistance. The data obtained are indicated in graphic form. There are 4 figures and 4 Soviet-bloc references.

ASSOCIATION: Institut fizicheskoy khimii AN SSSR (Institute of Physical Chemistry, AS USSR)

Card 2/4

S/080/61/034/009/CCB/016
D204/D305

AUTHORS: Solov'yeva, Z.A., Petrova, Yu.S., and Vagramyan, A.T.

TITLE: Effect of chromic acid concentration on the electrode
position of chromium

PERIODICAL: Zhurnal prikladnoy khimii, v. 34, no. 9, 1961.
2007 - 2012

TEXT: This is report II from a series of papers on investigating the rates of linked reactions in the electrodeposition of chromium. The effect of chromic acid concentration at constant and varying H_2SO_4 contents on the rate of deposition of metallic chromium, the rate of intermediate reduction of chromic acid and the rate of hydrogen evolution were studied. Also, the relationship between current distribution and CrO_3 concentration at constant and varying H_2SO_4 contents at various temperatures, and the variation of cathode potential with CrO_3 concentration at constant H_2SO_4 content

Card 1/3

S/080/61/034/009/008 006
D204/D305

Effect of chromic acid ...

were investigated. It was found that a decrease in chromic acid concentration increases the rate of chromium deposition sharply, the rate of increase depending on H_2SO_4 concentration, temperature and current density. The rate of intermediate reduction of chromic acid also increases and the rate of hydrogen evolution decreases. A decrease of chromic acid concentration with simultaneous increase in sulphuric acid concentration increases the rate of chromium deposition considerably. However, the rate of intermediate reduction of chromic acid is initially independent of the CrO_3 and H_2 L
 SO_4 concentration and subsequently decreases with a decrease in CrO_3 and H_2SO_4 concentration. The authors suggest that on changing the solution composition, the change in the rates of the linked reactions, taking place at the cathode during reduction of chromic acid in the presence of H_2SO_4 , is due to a change in composition and properties of the cathode film formed, from which discharge of Cr^{6+} ions occurs directly. There are 3 figures, 1 table and 4

Card 2/3

S/080/61/034/009/003/016
D204/D305

Effect of chromic acid ...

Soviet-bloc references.

SUBMITTED: August 29, 1960

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Card 3/3

S/080/62/035/008/005/009
D267/D308

AUTHORS: Solov'yeva, Z.A., Petrova, Yu.S., Klimasenko, N.L.,
and Vagramyan, A.T.

TITLE: Composition and properties of the cathode film forming
during the electrodeposition of chromium

PERIODICAL: Zhurnal prikladnoy khimii, v. 35, no. 8, 1962,
1806 - 1811

TEXT: The variation of concentrations of the ions Cr⁶⁺ and Cr³⁺
(volumetric method) and SO₄²⁻ (gravimetric method) was studied in
the deposited film, as a function of the concentration of H₂SO₄ and
H₂CrO₄ in the solution, and the variation of the rate of deposition
from H₂CrO₄ to Cr, in order to carry out a more complete investiga-
tion of the film composition. The coatings obtained in the course
of 2 min on a 6 cm² chromium-plated copper strip were either disso-
ved in distilled water or directly removed into a test tube. The so-
Card 1/2

Composition and properties of the ...

S/080/62/035/008/005/009
D267/D308

lutions contained 25, 50, 100 or 250 g/l H_2CrO_4 and 0.5 - 12.5 g/l H_2SO_4 , the temperature was $\sim 20^\circ C$ and the current density 250 or 500 ma/cm². Formation of two types of films was established: 1) Films obtained in the presence of H_2SO_4 distinguished by a macroscopic structure, soluble in the electrolyte in the absence of current and contributing to the reduction $H_2CrO_4 \rightarrow Cr$; 2) films obtained during an electrolysis without H_2SO_4 or when H_2CrO_4 acts without current on the cathode metal, possessing a microscopic structure, insoluble in the electrolyte. The latter do not contribute to the reduction of H_2CrO_4 . The relative concentration of Cr^{3+} and SO_4^{2-} in the films of the first type increases as the current density and the concentration of H_2SO_4 increases and as the concentration of H_2CrO_4 decreases. There are 4 tables.

SUBMITTED: June 22, 1961

Card 2/2

1D. 7400
5. 4600

AUTHORS:

Vagramyan, A. T., Krasovskiy, A. I., Petrova, Yu. S.
Solov'yeva, Z. A. (Moscow)

615
S/076/60/034/06/16/040
B015/B061

TITLE:

The Role of Passivation in the Electrodeposition of Metals

PERIODICAL: Zhurnal fizicheskoy khimii, 1960, Vol. 34, No. 6,
pp. 1255-1259

TEXT: The action of the rate of passivation on the electrochemical reduction of metal ions in aqueous solutions was examined. A series of experiments took place in manganosulfate solutions with and without additions of ammonium sulfate at pH = 8 and at 25°C. The results show (Table) that the reduction of the manganese²⁺ ions takes place through an activation of the electrode surface by ammonium sulfate. In a further series of tests a common electrolytic depositing of molybdenum and nickel from ammonium citrate solutions was examined. The rate of depositing of the nickel²⁺ rises through the activation of the surface with increasing concentrations of ammonia. One of the main obstacles to the reduction of metal ions is the passivation of the surface. Metals with a great

Card 1/2

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The role of Passivation in the
Electrodeposition of Metals

S/076/60/034/06/16/040
B015/B061

passivation tendency are hard to reduce on the cathode, and can be divided into three groups in this respect: 1) Metals which are deposited by relatively low overvoltage, such as Sn, Cd, Cu, Ag, etc., 2) Metals which are deposited at high overvoltage, such as Fe, Ni, Co, and Cr, Mn, etc., 3) Metals which cannot be deposited in pure form from aqueous electrolytes, such as Mo, W, U, Nb, Ti, Ta. In order to reduce metal ions, it is necessary to produce conditions which hinder passivation of the electrode surface, or at least strongly reduce it. R. I. Agladze is mentioned in the text. There are 4 figures, 1 table, and 4 references: 2 Soviet and 2 British.

ASSOCIATION: Akademiya nauk SSSR Institut fizicheskoy khimii Moskva
(Academy of Sciences of the USSR, Institute of Physical
Chemistry, Moscow)

SUBMITTED: August 6, 1958

Card 2/2

SOLOV'YEVA, Z.A.; VAGRAMYAN, A.T. (Moskva)

Oscillographic study of film formation on the cathode surface
in the electrolytic reduction of chromic acid. Zhur. fiz. khim.
36 no.4:752-756 Ap '62. (MIRA 15:6)

1. Akademiche nauk SSSR, Institut fizicheskoy khimii.
(Oscillography) (Chromic acid) (Electrolysis)

VAGRAMYAN, A.T., doktor tekhn. nauk, prof.; SOLOV'YEVA, Z.A., kand.
khimich. nauk

Electrolytic precipitation. Nauka i zhizn' 30 no.1:16-21
(MIRA 16:4)
Ja '63.

1. Zavedyushchiy laboratoriya elektroosazhdeniya metallov
Instituta fizicheskoy khimii AN SSSR.
(Electrometallurgy)

SOLOV'YEVA, Z.A.; VAGRAMYAN, A.T.

Effect of the cathode surface as affecting the character
of an electrochemical reaction. Dokl. AN SSSR 154 no.4:
918-921 F '64. (MIRA 17:3)

1. Institut fizicheskoy khimii AN SSSR.

L 46842-66 EWT(m)/EWP(t)/ETI IJP(c) JD/GD
ACC NR: AT6024972 (N) SOURCE CODE: UR/0000/65/000/000/0186/0189

32
B+1

AUTHOR: Solov'yeva, Z. A.

ORG: none

TITLE: Electrolytic preparation of iron-gold alloys

SOURCE: AN SSSR. Otdeleniye obshchey i tekhnicheskoy khimii. Zashchitnyye metalli-
cheskiye i oksidnyye pokrytiya, korroziya metallov i issledovaniya v oblasti elektro-
khimii (Protective metallic and oxide coatings, corrosion of metals, and studies in
electrochemistry). Moscow, Nauka, 1965, 186-189

TOPIC TAGS: iron alloy, gold alloy, electrodeposition, chemical composition

ABSTRACT: The electrochemical conditions of the codeposition of iron-gold alloys on a copper cathode at room temperature and the chemical composition of the deposits were investigated. The electrolyte was prepared by combining a cyanide solution of gold (0.5 g/l Au, 0.6 g/l KCN, 30 g/l Na₂HPO₄·12H₂O) and a citric acid solution of iron (40 g/l FeSO₄, 60 g/l citric acid) in various proportions. The composition of the deposits depends on the concentration of Au and Fe in the electrolyte and the current density. When the Fe:Au ratio in the electrolyte is 58 and the current density is about 30 mA/cm², the deposit contains approximately 85% gold. Such deposits, which constitute Fe-Au alloys, are insoluble in mineral acids, adhere well to copper and nickel bases, and have a grayish-yellow color which changes with the relative content

Card 1/2

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ACC NR: AT6024972

of the two metals. If the current distribution at the electrode is not uniform (as in the case of a lamellar cathode), separation of iron and gold in the deposit can be prevented by adding ammonia to the electrolyte. Addition of ammonia also decreases the gold content of the alloy: for example, when Fe:Au = 58 and at a current density of 30 mA/cm², the gold content drops from 85 to 10-20%. The current efficiencies in the presence of ammonia are 10-20%, and only 2-3% in its absence.

SUB CODE: 11,1307/SUBM DATE: 03Aug64/ ORIG REF: 002/ OTH REF: 004

Card 2/2 b1g

"APPROVED FOR RELEASE: 08/25/2000

CIA-RDP86-00513R001652330007-4

Effect of chlorine ions on cathodic polarization in chlorine
electrodeposition. Elektrokhimiia 1 no.5:451-456 by I.M.
(MIL 10.6)

Instytut Fizicheskoy Khimii AN SSSR.

APPROVED FOR RELEASE: 08/25/2000

CIA-RDP86-00513R001652330007-4"

SOVIET-EIA, L.V. V. N. M. K. T.

Effect of the angle of the cathode surface on the kinetics of
electrooxidation of chloro- α -D. Elektrokhimika 1 no. 2 p. 32.
(MRA 12-6)

1957-1965.

SOLOV'YEVA, Z.A.; VAGRAMYAN, A.T.

Cathodic passivation in chromate electrolytes with the additions
of chloride ions. Elektrokhimiia 1 no.4:413-417 Ap '65.
(MIRA 18:6)

1. Institut fizicheskoy khimii AN SSSR.

NIKITINA, A.A.; SOLOV'YEVA, Z.A.; SOMINSKAYA, Z.M.; VAGRAMYAN, A.T.

Mechanism of rhenium electrodeposition. Elektrokhimiia 1 no.6;
748-751 Je '65. (MIRA 18;7)

1. Institut fizicheskoy khimii AN SSSR.

SOLOV'IEVA, Z.A.

Electroreduction of chromic acid in the presence of fluoride
ion additions. Elektrokhimiia 1 no.7:812-817 Jl '65.
(MIRA 18:10)

1. Institut fizicheskoy khimii AN SSSR.

SOLOV'YEVA, Z.A., LAPSHINA, A.Ye.

Certain features of the electrodeposition of chromium from chromic acid solutions with fluoride ion additions. Elektrokhimia 1 no.8: 941-946 Ag '65.
(MIRA 18:9)

1. Institut fizicheskoy khimii AN SSSR.

SOLOV'YEVA, Z.A.

Effect of temperature on the electroreduction rate of chromic acid. Part 1. Elektrokhimiia 1 no.9:1088-1092 S '65.
(MIRA 18:10)

1. Institut fizicheskoy khimii AM SSSR.

SOLOV'YEVA, Z.A.

Effect of temperature on the electroreduction of chromic acid. Part 2. Elektrokhimiia 1 no.11:1366-1370 N '65.
(MIRA 18:11)

1. Institut fizicheskoy khimii AN SSSR.

SOLOV'YEVA, Z.I. (Donetsk, ul. Gor'kogo, d.25)

Apparatotherapy in spinal paralysis of traumatic origin. Ortop.,
travm.i protez. no.4:32-35 '62. (MIRA 15:5)

1. Iz Donetskogo instituta travmatologii, ortopedii i protesirovaniya (dir. - kand.med.nauk T.A. Revenko) i Donetskogo proteznogo zavoda (dir. - L.P. Polotskiy).
(SPINE—WOUNDS AND INJURIES) (PROSTHESIS)
(PARAPLEGIA)

AUTHORS: Perfilov, N. A., Solov'yeva, Z. I. SOV/89-5-2-12/36

TITLE: On a Complex Nuclear Fission of Uranium Under the Influence of Neutrons of an Energy of 14 MeV (O slozhnom delenii yader urana pod deystviyem neytronov s energiyey 14 Mev)

PERIODICAL: Atomnaya energiya, 1958, Vol. 5, Nr 2, pp. 175-176 (USSR)

ABSTRACT: In various papers (Refs 1-3) (with thick photoemulsions as well as with ionization chambers) the occurrence of a far-reaching α -particle was observed when the uranium nucleus undergoes fission. This α -particle is formed during the moment of fission. The angle between the direction of flight of this α -particle and the light fission fragments is about 80° . A maximum was observed within the domain of 16-17 MeV in the energy distribution of the α -particles. By means of nuclear photoplates saturated with uranium salt (exposed to a neutron beam of 14 MeV-neutrons) fission of the uranium was investigated with the emission of a far-reaching α -particle. Eight cases of a fission with emission of a far-reaching α -particle were found among 10 000 normal fission processes. From these and earlier measurements the conclusion may be drawn that under the influence of fast neutrons fission of the uranium

Card 1/2

On a Complex Nuclear Fission of Uranium Under the
Influence of Neutrons of an Energy of 14 MeV

SOV/89-5-2-12/36

nucleus with 3 fission fragments corresponds to about 1 000 to
1 300 cases of normal fission. This process, therefore, is about
3 to 4 times less probable than in the case of fission of U²³⁵
by thermal neutrons. There are 2 figures and 4 references, 1 of
which is Soviet.

SUBMITTED: February 10, 1958

Card 2/2

Sov. Lit. Obj. VD,
1.3
PHASE I BOOK EXPLOITATION

SOV/3503

Akademiya nauk SSSR. Radiyevyy institut

Trudy, t. IX (Transactions of the Radium Institute, Academy of Sciences USSR,
Vol. 9) Moscow, Izd-vo AN SSSR, 1959. 287 p. Errata slip inserted.
1,700 copies printed.

Ed.: N.A. Perfilov, Doctor of Physical and Mathematical Sciences; Ed. of Publishing
House: G.M. Aron; Tech. Ed.: A.V. Smirnova.

PURPOSE: The volume is intended for physicists.

COVERAGE: The book represents volume 9 of the Transactions of the Radium Institute
and contains the results of studies conducted at the Institute chiefly from
1955 to 1956. There are a number of articles dealing with the study of nuclear
reactions occurring with particles of different energies ranging from several
eV up to hundreds of MeV. Others treat different problems of the physics of
neutrons. Results of studies of various neutron sources, neutron energy distri-
bution in a moderator (water), and other problems connected with the theory of
neutron interaction with matter are presented. The majority of the articles

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Transactions of the Radium (Cont.)

SOV/3503

are concerned with problems of method. The authors provide a complete description of the construction of equipment and of the results of tests performed under laboratory conditions. No personalities are mentioned. References accompany individual articles.

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Card 1/8

PERFILOV, N.A.; SOLOV'YEVA, Z.I.

Special cases of the fission of uranium nuclei induced by slow
neutrons. Trudy Radiev, inst. AN SSSR 9:5-40 '59. (MIRA 14:6)
(Uránium) (Nuclear fission)

PERFILOV, N.A.; SOLOV'Yeva, Z.I.

Angular distribution of long-range α -particles associated
with the fission process. Zhur.eksp.i teor.fiz. 37 no.4:
1157-1159 O '59. (MIRA 13:5)
(Alpha rays) (Fission)

Complex Fission of Uranium Nuclei by 2.5-mev
Neutrons. Letter to the Editor

77245
SOV/89-8-2-10/30

double events, i.e., the branching ratio is 1:600. Comparing it to the 1:350 ratio for thermal neutrons and 1:1,100 ratio for 14-mev neutrons, the author concluded that the probability decreases with the increase of neutron energy. In the thermal energy region the fission is mostly due to U²³⁵. In the present study with 2.5-mev neutrons, the author performed a second experiment with a uranium salt containing five times less U²³⁵ isotope than the natural mixture. There was no noticeable change in the probability of triple fission, and the author concluded that he observed triple fission due to U²³⁸ and not U²³⁵. Other aspects of the event were similar to those at other neutron energies. The fission was asymmetric; mean ratio of the ranges of the light and heavy fragments $R_L/R_H = 1.3$. The α -fragment angular distribution with respect to the direction of the light fragment is given in Fig. 1. Figure 2 shows the α -particle range distribution. N. A. Perfilov showed interest

Card 2/4

Complex Fission of Uranium Nuclei by 2.5-mev
Neutrons. Letter to the Editor

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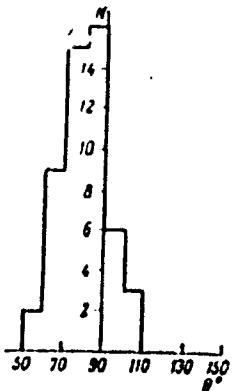


Fig. 1. Angular distribution of alpha-particles with respect to direction of flight of the light fragment.

Card 3/4

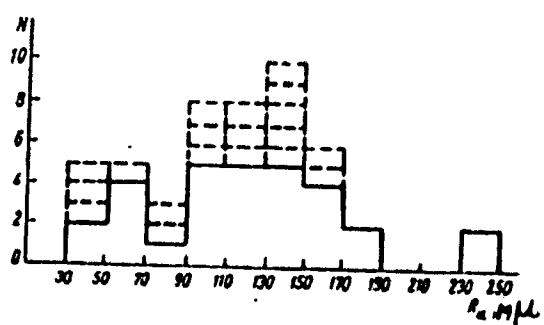


Fig. 2. Distribution of alpha-particle ranges during uranium fission due to action of 2.5-mev neutrons:
--- tracks which went out of the photolayer.

Complex Fission of Uranium Nuclei by 2.5-mev
Neutrons. Letter to the Editor

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in the work. There are 2 figures; and 3 references,
1 Soviet, 1 French, 1 U.S. The U.S. reference is:
E. Titterton, Phys. Rev., 83, 673 (1951).

SUBMITTED: June 20, 1959

Card 4/4

Selby 4/24/64

21-5280

34-6000

ATTACH:

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FROM:

INVESTIGATION OF AD AND MR PROJECT ON PROTON SOURCE OF

PROTONS

REMARKS:

Partial description of investigation results, 1960.

Vol. No. 1, pp. 245 - 300

TEXT: The authors of the paper under review offer the first results obtained from their investigation of AD and MR fragmentation (nuclear disintegration) in multicellular particles with T = 0 by proton irradiation of small amounts of substances containing one isotope of each of the elements (100% natural), namely benzene, cyclohexane, the isobutane (isobutyl bromide), propyl bromide, and a vinyl phenyl ether. The additional papers were numbered by V. N. Slobodin and co-workers also. In his interpretation of the results, each molecule also. Isotropisms were selected as constant traits of particles with T = 0.

Card 1/1

Card 2/1

Presentation of the results of proton scattering

Scanning electron microscopy

of the metal insulator interface after sputtering. Fig. 2 illustrates the fragmentation cross section as a function of E_p . In the range of proton energies around 1 MeV there appears a steep rise in the cross section, a multiplicity of fragments produced. The quantity of atoms with two or more tracks of multicharged particles is found to increase with the energy of bombarding protons.

Usually about 10% of 100 MeV O₂ ions have a double ionization. The number of fragments decreases in a practically linear manner with increasing energy. Below the charge distribution, a distribution curve from the one found at the charge distribution of angular distribution is shown. The angular distribution of the bombarding particles was measured by means of the fragments. Their angular distribution is shown in Fig. 3. It is found that the angular distribution of fragments is in agreement with the previous distribution, i.e., more frequent at direct impact, with less frequent and with more frequent diffraction. The angular distributions were analyzed by an interesting fragment counter. The angular distribution ratio is 1.1 at 100 MeV. The angular distribution has been calculated by the method of moments with respect to the given three parameters:

Card 1/4

CONT'D

DATE REC'D.: AUGUST 11, 1986

REF ID: A6513R001652330007-4

S/053/60/071/03/03/008
B006/B063

AUTHORS: Perfilov, N. A., Romanov, Yu. F., Solov'yeva, Z. I.
TITLE: Fission of Heavy Nuclei With Emission of Long-range α -Particles /⁹
PERIODICAL: Uspekhi fizicheskikh nauk, 1960, Vol. 71, No. 3, pp. 471-483

TEXT: Long range particles are said to be formed in a nuclear fission such as is shown in Fig. 1, where three charged particles are emitted. The nature of the long-range component is discussed first. Then experiments are described by which Z^2/m and mZ^2 of these particles were determined leading to their identification as α -particles. The probability of such a complicated fission event, as well as the methods used to determine the relative probability of triple fissions are discussed next. Photoemulsions and U^{233} or Pu^{239} are most frequently used. The authors give a brief description of the method of the double ionization chamber (Ref. 6) and of the possibility of determining the relative probability by integrating the α -particle distribution function (Ref. 10). Then, the ✓

Card 1/3

Fission of Heavy Nuclei With Emission of
Long-range α -Particles

S/053/60/071/03/03/008
B006/B063

apparatus used in Ref. 9 is described (Fig. 2), and a comparison is made between the results obtained by various methods for U^{235} fission induced by thermal-neutrons (Table 1). Reference is made to numerous Western authors, whose results are compared with one another and discussed. The next part of the present paper deals with the energy distribution of long-range α -particles. The spectrum may be analyzed by means of photographic plates, absorption in thin aluminum foils, an ionization chamber with a grid, and the deviation in the magnetic field. Methods described in some Western articles are discussed in more detail, and distribution curves obtained by several authors are compared with one another in Fig. 3. Good agreement was found between the data obtained by various methods. In this connection the authors briefly refer to their own investigations (Refs. 15 and 16). The fourth section presents details on methods and results of the determination of angular distribution. Fig. 4 shows a typical distribution curve. The fifth section deals with the energy spectrum of the fragments of a triple fission. Two peaks were found for U^{235} , one at 10 Mev and another at 7 Mev. Investigations by V. N. Dmitriev, L. V. Drapchinskiy.

Card 2/3

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Fission of Heavy Nuclei With Emission of
Long-range α -Particles

S/053/60/071/03/03/008
B006/B063

V. A. Petrzhak and Yu. F. Romanov (Ref. 25, Fig. 5), as well as by
N. Mostovoy, T. A. Mostovaya, M. Sovinskiy, and Yu. S. Saltykov (Ref. 26)
are discussed in this connection. Fig. 6 shows a comparison between the
results of Refs. 9 and 25. Some hypotheses on the mechanism of triple
fission, published in Western articles, are discussed in the final part.
There are 7 figures, 1 table, and 31 references: 10 Soviet, 13 American,
5 French, and 4 British.

Card 3/3

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PERFILOV, N.A.; SOLOV'YEVA, Z.I.; FILOV, R.A.

Triple fission of uranium nuclei by fast neutrons. Zhur.eksp.i teor.
(MIRA 14:7)
fiz. 41 no.1:11-12 J1 '61.

1. Radiyevyy institut AN SSSR.
(Nuclear fission) (Uranium isotopes) (Neutrons)

PERFILOV, N....; SOKOV'YAN, T.I.; FILOV, . .A.; KULAKHOV, G.I.

Spontaneous triple fission of curium-242. Dokl. AN SSSR 136
no. 3:581-582 Ja '61. (KIRD 14:2)

1. Radijevyy institut imeni V.G. Khlopin A.N. SSSR. Predstavleno
akademiku D.N. Konstantinovym.
(Curium-decay)

L1121
S/056/62/043/004/004/061
B102/B186

24660
AUTHORS:

Bolovtseva, Z. I., Filov, R. A.

TITLE:

Dependence of the kinetic energy of fragments on the α -particle energy during triple fission of uranium

PERIODICAL: Zhurnal eksperimental'noy i teoreticheskoy fiziki, v. 43,
no. 4(10), 1962, 1146 - 1148 .

TEXT: Studies of the energy distribution of U^{235} fragments formed in double and triple fission showed that, the most probable total kinetic energy of double-fission fragments, equals the energy triple-fission fragments plus the probable energy of the long-range α -particle. Further studies were made to find out whether this relation holds generally. π -80 (P-80) nuclear emulsion plates were impregnated with uranium salt and irradiated with thermal neutrons from a reactor. α -particle and total fragment energies were determined from fission ranges, and the energy distributions were established for the intervals 6-14 Mev (10.4 Mev mean value), 14-17 Mev and 17-30 Mev (21.0 Mev mean value). The most probable fragment ranges were found to be shifted by ~10 Mev in dependence on the mean energy X

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Dependence of the kinetic energy ...

S/356/62/043/004/004/061
B102/3186

of the long-range α -particles. This confirms that the relation stated above is valid also for energies other than the most probable. The triple fission mechanism is discussed in the context of a dumbbell-shaped nuclear model. There is 1 figure.

SUBMITTED: April 14, 1962

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L 10677-63

EPF(n)-2/S/T(m)/BDS--FFTC/ASD/ FNL/SSD--Pu-4

ACCESSION NR: AP3002265

8/0089/63/014/ 006/0575/0577
60AUTHOR: Perfilov, N. A.; Solov'yeva, Z. I.; Filov, R. A.TITLE: Triple fission of U sup 235 by neutrons of 14 mev energySOURCE: Atomnaya energiya, v. 14, no. 6, 1963, 575-577

TOPIC TAGS: triple uranium fission, neutron uranium fission

ABSTRACT: The triple splitting of uranium nucleus (fission plus the formation of a high energy alpha particle) was observed in a previous work to be more probable for bombardment with thermal neutrons than with fast ones. However, these observations were made with different isotopes: thermal neutrons with U sup 235, fast neutrons with U sup 238. Therefore, neutrons of 14 Mev were used with U sup 235 in this study. Photoemulsions P-9-0 were soaked in uranium salt solutions and irradiated in the neutron beam from the reactor $t(\alpha, n)$ He sup 4. About 100 thousand fission events were observed, among them 65 triple ones. The energy spectrum of alpha particles and their angular distribution were observed and given in diagrams. They are approximately the same for thermal and for fast neutrons. The probability of triple splitting decreases with the increase of the isotopic mass.

Orig. art. has: 3 figures.

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L 13613-63

EWT(m)/BDS AFFTC/ASD

ACCESSION NR: AP3003107

S/0056/63/044/006/1832/1836 54
53

AUTHOR: Perfilov, N. A.; Solov'yeva, Z. I.; Filov, R. A.; Khlebnikov, G. I.

TITLE: Ternary fission of plutonium 19

SOURCE: Zhurnal eksper. i teor. fiziki, v. 44, no. 6, 1963, 1832-1836

TOPIC TAGS: ternary fission of plutonium, Alpha particle energy spectra, plutonium thermal fission, uranium complex fission

ABSTRACT: The energy spectra of long-range Alpha particles produced in the spontaneous fission of Pu²³⁸ and Pu²⁴⁰ and in thermal fission of Pu²³⁹ have been studied by the nuclear emulsion method with an aim at comparing both the fission probabilities and the fission Alpha-particle energy spectra of the different isotopes. Electrolytic films of Pu²³⁸ and sup 240, containing 78 plus or minus 4 and 450 plus or minus 25 microgram respectively were used in the spontaneous fission test, and Pu²³⁹ film irradiated with neutrons from the reactor of FTI AN SSSR was used to obtain the alpha-particle energy spectrum from thermal-neutron fission. The photographic plates were scanned with a microscope and the resultant histograms were tested for fits to Gaussian distributions with various maxima and half-widths. The spectrum shapes are discussed and compared

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ACCESSION NR: AP3003107

with the results for complex uranium fission. Logical reasons for representing the Alpha-particle spectra as Gaussian or near-Gaussian distributions are advanced and agreement in the case of ternary fission of Pu sup 240 is noted with recent work by R. A. Nobles (Phys. Rev. v. 126, 1508 (1962)). "In conclusion, the authors wish to thank V. M. Kulakov for experimental assistance." Orig. art. has: 2 figures, 1 formula, and 2 tables.

ASSOCIATION: none

SUBMITTED: 17Jan63 DATE ACQ: 23Jul63 ENCL: 003

SUB CODE: 00 NO REF Sov: 008 OTHER: 007

Card 2/52

ACCESSION NR: AP4042591

S/0056/64/046/006/2244/2245

AUTHORS: Perfilov, N. A.; Solov'yeva, Z. I.; Filov, R. A.

TITLE: Alpha particle spectrum of ternary spontaneous fission of
Cm-244

SOURCE: Zh. eksper. i teor. fiz.. v. 46, no. 6, 1964, 2244-2245

TOPIC TAGS: curium, alpha particle, fission product, energy dis-
tribution, alpha spectroscopyABSTRACT: The investigation was undertaken in order to check whether
the spectral characteristic of the fission alpha particles is indeed
independent of the mass number, as was observed by the authors
earlier (Atomn. energ. v. 14, 575, 1963; ZhETF v. 44, 1832, 1963).
The test was also aimed at checking the correctness of the value ob-
tained for the most probable energy of the fission alpha particles
for Cm²⁴². The same experimental setup was used as in the earlier

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ACCESSION NR: AP4042591

investigation, and geometrical corrections were introduced into the experimentally observed energy distribution. The obtained energy spectrum for the ternary fission of Cm²⁴⁴ was found to have a maximum near 15.5 ± 0.5 MeV and, assuming a Gaussian distribution, a width 11.5 ± 0.5 MeV at half height. The values obtained for Cm²⁴² are $15.5 \pm$ and 12 ± 1 , respectively, indicating that the spectral characteristics agree, within the experimental error, also for two curium isotopes, as they did agree before in the case of U and Pu. However, the shift in the most probable energy of ternary-fission alpha particles cannot be established on the basis of data on three elements only. "The authors thank A. S. Krivokhatskiy for assistance." Orig. art. has: 1 figure.

ASSOCIATION: None

SUBMITTED: 01Apr64

ENCL: 01

SUB CODE: NP

NR REF SOV: 004

OTHER: 003

Card 2/3

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CIA-RDP86-00513R001652330007-4

PERFILOV, N.A., SOLOV'YEEVA, Z.I.; FILOV, R.A.

Spectrum and particles in the triple spontaneous fission of
cm₂₄₄. Zhur. eksp. i teor. fiz. 46 no.6:2244-2245 Ja '64.
(MIFI A 17:10)

APPROVED FOR RELEASE: 08/25/2000

CIA-RDP86-00513R001652330007-4"

DORMAN, L.I.; INOZEMTSEVA, O.I.; MAZARYUK, Ye.A.; SOLOV'YEVA, Z.I.

Modulation of the solar diurnal effect and the possibility for
establishing variations in cosmic ray intensity as measured
in sidereal time. Izv. AN SSSR. Ser. fiz. 29 no.10:1898-1901 0
'65. (MIRA 18:10)

SOLOV'YEVA, Z. I.

Nov/Dec 47

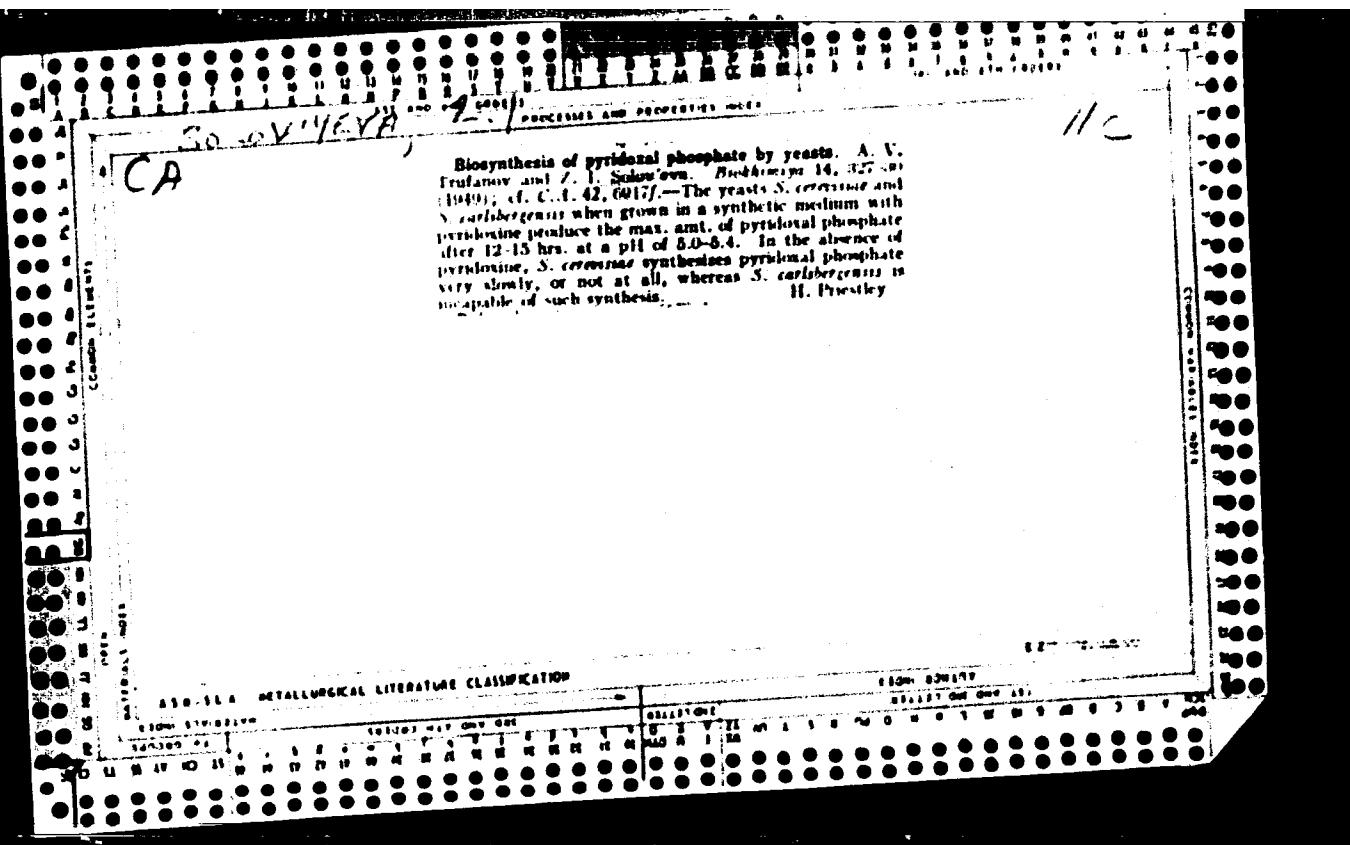
USSR/Medicine - Pyridoxal
Phosphate
Medicine - Biochemistry

"Pyridoxal Phosphate Synthesis by Animal Tissues," A. V.
Trufanov, V. A. Kiranova, Z. I. Solov'yeva, Lab of Chem of Vitamins,
Nutrition Inst, Acad Med Sci USSR, 8 pp

"Biokhim" Vol XII, No 6

Explained synthesis of pyridoxal phosphate (codecarboxylase) in vitro
in sections of livers, kidneys, heart, muscles and brains of normal rats
in the presence of pyridoxene. Synthesis of pyridoxal phosphate is
related to the enzymatic system of the cells which have optimum pH of
about 7.2. Maximum synthesis occurs when pyridoxene and fresh tissues are
mixed in a ratio of 2.5 mg doxane, to 1 g of tissue. Submitted 18 Feb 48.

62/49T50



SOLOV'YEVA, Z. I.

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Absorption, elimination, and distribution of radioactive thiopental in rat organs and tissues. L. I. Gribenauk and Z. I. Solov'yeva. *Farmakol. i Toksikol.* 17, No. 1, 22-8 (1964). When Na thiopental (contg. S³⁵) was given to rats, 61.61% was eliminated in 4 days (mostly in the first 2 days), 40.18% appeared in oxidized form and 9.30% as unchanged Na thiopental or its primary degradation products. Nephritis due to CCl₄ inhibits elimination of Na thiopental and its degradation products by the kidneys. J. V. S.

GREBENNIK, L.I.; SOLOV'YEVA, Z.I.

Absorption, excretion, and distribution of radioactive thiopental sodium in the organs and tissues of rats. Khim.i med. no.11:104-110 '59. (MIRA 13:6)

(THIOPENTAL)

GORYACHEVA, N.S.; SOLOV'YEVA, Z.I.

Quantitative determination of cutisone in preparation and in
tablets. Med.prom, 16 no.4:33-36 Ap '62. (MIRA 15:8)

1. Vsesoyuznyy nauchno-issledovatel'skiy khimiko-farmatsevti-
cheskiy institut imeni S.Ordzhonikidze.
(BENZALDEHYDE)

SOLOV'YEVA, Z.M.

Some characteristics of the performance of the equipment for
casting and vulcanization of diaphragms for molder-vulcanizers
in the Dnepropetrovsk Tire Plant. Kauch.i rez. 22 no.2:43-46
(MIRA 16:2)
F '63.

1. Dnepropetrovskiy shinnyy zavod.
(Dnepropetrovsk—Rubber industry—Equipment and supplies)
(Vulcanization)

SOLOV'YEVA, Z.M.

Studying the effect of the design features on the operative properties of the membranes of molder-vulcanizers. Kauch.
i rez. 22 no.8:40-46 Ag '63. (MIRA 16:10)

1. Dnepropetrovskiy shinnyy zavod.

SOLOV'YEVA, Z.M., ALEKSANDROV, V.V.

Technological process of the manufacture of diaphragms for
shaper-vulcanizers used by the Dnepropetrovsk Tire Plant.
Kauch. i rez. 24 no.8, 35-41 '65. (MIRA 18:10)

1. Dnepropetrovskiy shinnyy zavod.

STEPANOV, V.N., prof., doktor sel'skokhoz. nauk; GOLININA, N.E., nauchnyy
sotrudnik; SOKOLOV'YEVA, T.M., nauchnyy sotrudnik

Yield capacity and productivity of farm crops during crop rotations
of various combinations in central regions of the non-Chernozem belt.
(MIRA 18:5)
Izv. TSKHA no.5:106-121 '64.

1. Kafedra rasteniyevodstva Moskovskoy ordena L'nina sel'skokho-
zyaistvennoy akademii imeni Timiryazeva.

LYUBCHENKO, A.P., MASHAROV, N.V., SHERMAN, D.G., SOLOV'YEVA, Z.P.

Microdistribution of elements in cast iron altering the face
cut of graphite crystals. Fiz. met. i metalloved. 13 no.4 p.11
572 O '64. (Mir 18:4)

1. Khar'kovskiy zavod transportnogo mashinostroyeniya imeni
Malyshova.